Amendments to the Claims:

This listing of claims will replace all prior versions and listings of claims in the application.

Listing of Claims:

1-15. (Canceled)

16. (Currently Amended) A pharmaceutical composition comprising;
an extended-release first portion made of one or several units containing therapeutically effective amounts of NSAIDs mixed with at least one retardant material for extended release delivery of the non-steroidal anti-inflammatory drugs (NSAIDs) presenting a controlled availability of the non-steroidal anti-inflammatory drug (NSAIDs) alongside the gastrointestinal tract;
an immediate release second portion made of a powder of one or several units containing therapeutically effective amounts of the stabilised gastroprotective prostaglandin and a pharmaceutical carrier for the immediate release of said stabilised gastroprotective prostaglandin, and
wherein the extended release first portion and the immediate release second portion are encapsulated within a capsule made of hydroxyl-propyl-methyl-cellulose (HPMC) polymer.

17. (Canceled)

18. (New) The pharmaceutical composition according to claim 16, wherein the first and second portions are separated by a third portion.

19. (New) The pharmaceutical composition according claim 16 , wherein the non- steroidal anti-inflammatory drug (NSAID) is selected from the group consisting of aceclofenac, diclofenac, difflunisal, fenbufen, flufenamic acid, ibuprofen, indomethacin, ketoprofen, meclofenamate sodium, meloxicam, mefenamic acid, nabumetone, naproxen, piroxicam, suprofen, tiaprofenic acid, acetylsalicylic acid, flurbiprofen, ketorolac, oxaprozin, sulindac, tenoxicam, tiaprofenic acid and suitable salts thereof.

20. (New) The pharmaceutical cornposition.according to claim 16, wherein the retardant material of the first portion is selected from the group consisting of lipidic materials, acrylic and methacrylic acid polymers and copolymers, alkyl celluloses, gums, protein derived materials and a mixture thereof.

21. (New) The pharmaceutical composition according to claim 16 , wherein the prostaglandin is a "E-series' prostaglandin selected from the group consisting of PGEl , PGE2, misoprostol, enoprostol, enisoprost, rosaprostol, and miraprostol.

22. (New) The pharmaceutical composition according to the claim 21, wherein the gastroprotective prostaglandin is misoprostol stabilized by a dispersion in hydroxy-propylrnethylcellulose (HPMC) or polyvinylpyrrolidone (PVP).

23. (New) The pharmaceutical composition according to claim 16, wherein the stabilized gastroprotective prostaglandin is stabilized misoprostol and the non-steroidal anti-inflammatory drug (NSAID) is selected from the group consisting of diclofenac, ketoprofen, ibuprofen, meloxicam, and naproxen.

24. (New) A method for the treatment of inflammatory conditions or diseases in a mammal patient, including the human, that comprises the step of administrating a sufficient amount of the pharmaceutical composition according to claim 16, to said mammal patient.

25. (New) The method according to claim 24, wherein said inflammatory condition or disease is osteoarthitis or rheumatoid arthritis.

26. (New) The method of claim 24, wherein the pharmaceutical composition is administrated as a dual release formulation allowing a one a day or twice a day dosing into humans.

27. (New) The method of the claim 20, wherein the lipidic material is selected from the group consisting of waxes, glycerides or aliphatic alcohols.

28. (New) The method of claim 20, wherein the acrylic or methacylic acid polymer is selected from the group consisting of methylacrylate polymers, methyl methylacrylate copolymers, ethoxyethyl methacrylate polymers, cyanoethylmethacrylate polymers, aminoalkyl methacrylate copolymers, poly (acetylic acid) , poly (methacrylic acid), methacrylic acid alkylamine copolymers, poly(methyl methacrylate) polymers, poly (methacrylic acid)(anhydride), polymethacrylate polymers, polyacrylamide, poly (methacrylic acid anhydride), glycidyl methacrylate copolymers or a mixture thereof.

29. (New) The method of claim 20, where the alkyl celluloses are selected from the group consisting of hydroxypropylmethylcelluloses (HPMC), hydroxyethylcelluloses (HEC), hydroxypropylethylcelluloses (HPEC), hydroxypropylcelluloses (HPC), methylcelluloses (MC), ethylcelluloses (EC), sodium carboxymethylcelluloses (NaCMC), and a mixture thereof.